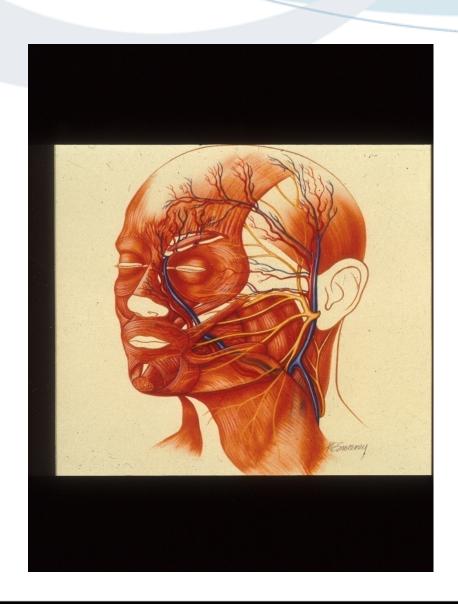
Electroneuronography (ENoG)





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ENoG: Presentation Overview

- **☐** Etiologies
- ☐ Alternative facial nerve tests
- ☐ Facial nerve anatomy
- ☐ Facial nerve physiology
- □ Bell's Palsy
- ☐ H-B Facial Grading System
- □ Stimulation Techniques
- **☐** Recording Techniques
- ☐ Interpretation & reporting guidelines.



Facial Nerve Paralysis

Facial nerve paralysis is absolutely debilitating.

Primary debilitation involves emotional and psychological impact of facial disfiguration. Socialization and community participation is extraordinarily limited and difficult for many of these patients.

Secondary debilitation involves physical limitations, difficulties speaking, drinking, eating and facial expression.



Facial Nerve Disorders: Possible Etiologies

- **□** Mumps
- □ Chicken-pox
- **☐** Gilllain-Barre syndrome
- ☐ Central nervous system disorders (i.e., stroke)
- **□** Glomus jugulare
- □ Meningioma
- ☐ Facial nerve neuroma
- □ Trauma



Facial Nerve Disorders: Etiologies cont'd...

- □ Bell's Palsy (more on this in a moment)
- □ latrogenic (surgically induced)
- ☐ Temporal bone trauma secondary to MVA
- □ Otitis media
- ☐ Herpes zoster oticus
- □ Multiple sclerosis
- Melkersson-Rosenthal syndrome
- Mastoiditis



Other facial nerve tests:

- ☐ Hilger test
- □ Electromyography (polyphasic APs…)
- □ Acoustic reflex testing
- **□** Antidromic nerve potentials
- MRI and CT
- Maximal/minimal nerve stimulation tests
- ☐ Transcranial magnetic stimulation
- ☐ Blink reflex tests
- □ Others....



The Essence of ENoG....

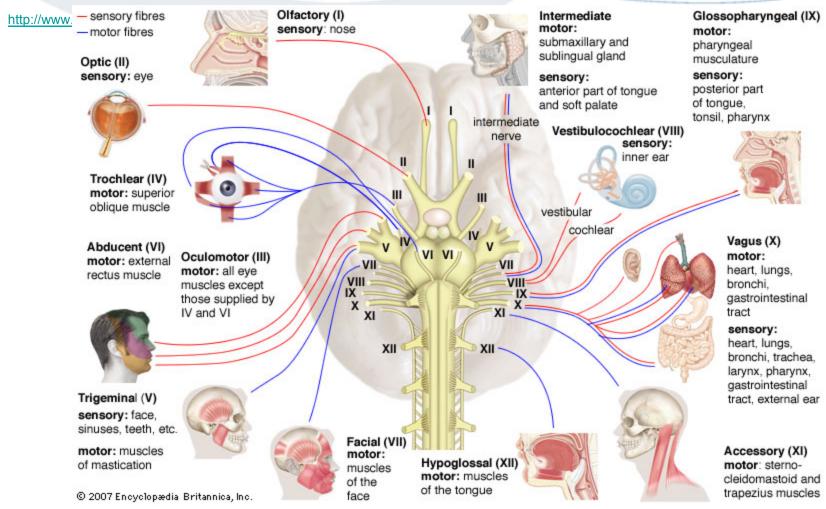
ENoG is the electrical stimulation of the facial nerve at (or near) the stylomastoid foramen (SMF) with measurements taken from the nasio-labial fold.

ENoG is the only "relatively objective" measure of facial nerve integrity.

ENoG compares the normal side to the abnormal side to quantify and prognosticate.

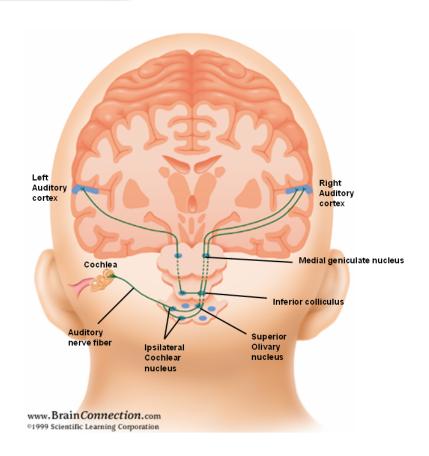


Cranial nerve innervation

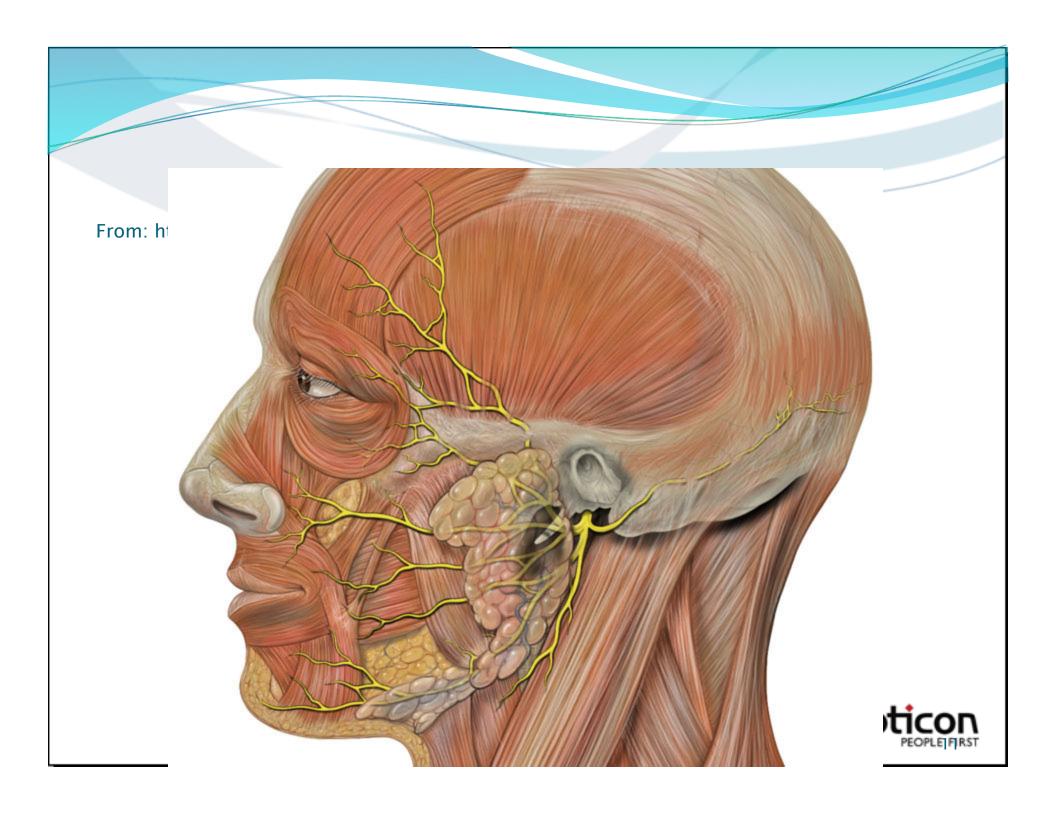




VII & VIII travel together...from brainstem to distal end of the IAC...







Facial Nerve Anatomy...

Facial Nerve (cn VII) has 10,000 fibers (2/3rds motor, 1/3rd sensory)

Normal facial motion requires only half the motor fibers.



Facial Nerve Anatomy ...

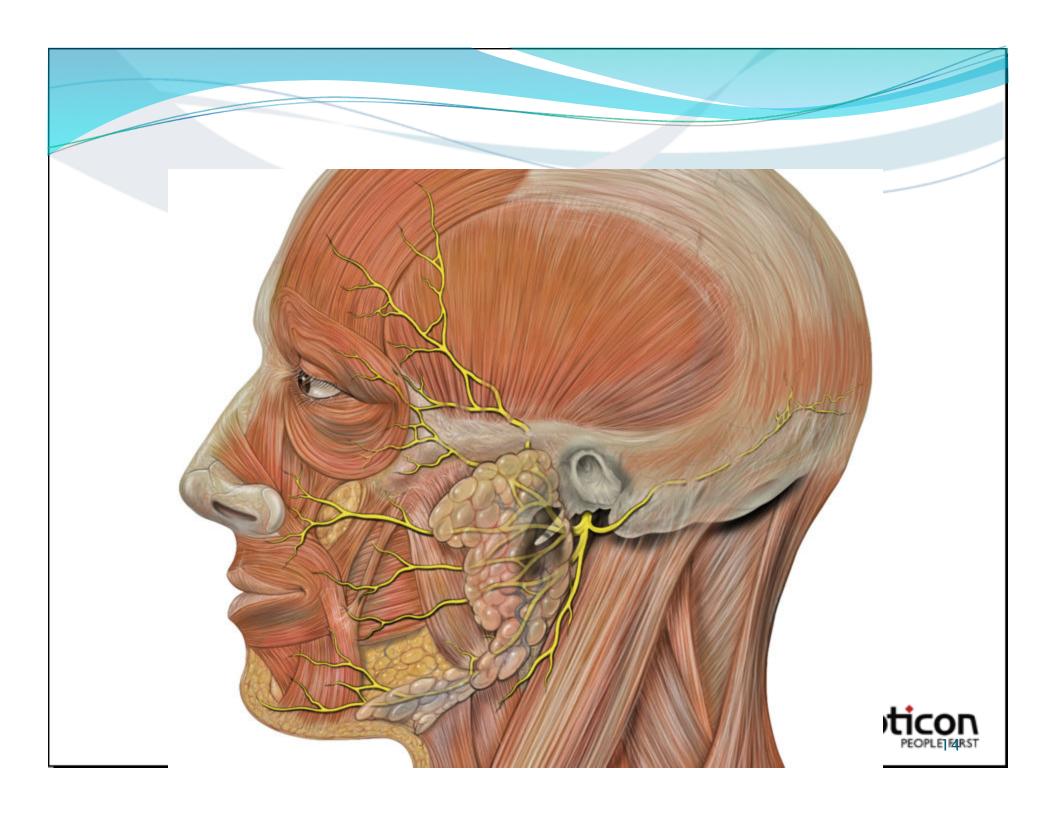
Facial nerve has five landmark segments:

- 1) labyrinthine
- 2) tympanic
- 3) pyramida
- 4) mastoid
- 5) stylomastoid foramen (SMF)

The facial nerve emerges from SMF, enters parotid gland and divides:

- 1) Superior division (temporal, zygomatic, buccal)
- 2) Inferior division (buccal, mandibular, cervical)





3 Types of Facial Nerve Injury:

Neuropraxia: Most common finding with BP. Paralysis without

peripheral nerve degeneration. ENoG is normal or

reduced response, nerve fibers and sheath are

intact, but not responsive to volitional commands.

Neurotmesis: Worst possible outcome. ENoG no response

(i.e. flat line). Total anatomic separation,

very poor prognosis

Axnotmesis: Inner nerve fiber disruption with intact outer

casing (epineurium). ENoG no response (i.e., flat line).

(ENoG cannot differentiate neurotmesis from axonotmesis)



Bell's Palsy Overview:

Facial muscles weak or non-responsive.

Caused by injury to the 7th cranial nerve.

Sir Charles Bell, Scotland, 200 years ago.

Worldwide incidence 0.02%:

1 in 5000 people worldwide, 40,000 in the USA annually.

Left/right, Male/female, race, L and R sides equal.

Older people more likely, children are possible.

Diabetics four times more likely than non-diabetes

Last trimester of pregnancy higher risk for BP.

HIV & Sarcoidosis increase the odds of acquiring BP (Sarcoidosis is an inflammation, produces microscopic granulomas)



BP Overview cont'd...

Both sides simultaneously, rare (less than 1 %)

Does not affect other body systems.

BP is often noticed first thing in the AM...

Early symptoms (dry eye, tingling mouth/lips, weak mouth (slurping). May take a day or two to fully engage.

BP usually "peaks" within 72 to 96 hours.

BP is not contagious.

Recovery: 50% complete recovery in 4 to 8 weeks.

35% recover in less than a year.

Recurrence: 8 percent recur, may take ten years until 2nd event.



BP Treatment Options:

- □ Surgical intervention
- □ Wait and watch approaches
- Medical management
 - ✓ Acyclovir
 - ✓ Steroids...
- ☐ Eye care:
 - ✓ Patch
 - ✓ Tape
 - √ Shades
 - ✓ Artificial tears....



House-Brackmann (HB) Facial Grading Scale:

HB scale approximates quantity of volitional motion based on clinical facial presentation.

HB grossly describes characteristics and degree of facial nerve motion using subjective analysis.

HB scale has six grades, each grade is reported as a fraction.



House-Brackmann Facial Grading

Grade one (1/6): normal

Grade two (2/6): slight/mild weakness

Grade three (3/6): moderate weakness with normal eye closure

Grade four (4/6): moderate weakness without eye closure

Grade five (5/6): severe weakness

Grade six (6/6): total facial paralysis



When to test? Which question are we answering....

Consider, only grade six (6/6) presentations *require*EnoG testing, if the purpose is to determine whether the facial nerve is intact. If we're "tracking" the progress of function or dysfunction, ENoG can be used to monitor progress.

Difficult to discern 5/6 from 6/6 due to masseter or contralateral pull



Wallerian Degeneration:

Wallerian Degeneration (WD) takes approximately 72 hrs for denervation to completely impact neural fibers.

If you test HB 6/6 facial one hour post-onset BP, the likely result is a normal ENoG because neural fibers are physiologically intact, although non-functional volitionally, yields "false negative" test.

Wait 72 hours before first ENoG to allow complete WD.

WD occurs proximal to distal to damage in CNS and PNS.



Timing the ENoG Test:

ENoG "valid" from 3 to 21 days.

1st test 72 hrs post-onset, retest 3 to 5 day intervals.

No guidelines after 21 days....



Surgical Management of BP Laryngoscope 109(8), August 1999, Gantz, Rubinstein, Gidley & Woodworth

54 subjects < 90% denervation all returned to HB grade I or II within 7 months post-onset.

19 subjects >90% denervation, with no f.n. decompression 58% chance of outcome HB grade III or IV at 7 mos post-onset.

31 subjects >90% denervation, with f.n. decompression 91% chance of HB grade I or II.



ENoG:Stimulus Parameters

Transducer: Pair of electrodes

Site: Stylomastoid foramen

Orientation: anode (+) anterior ("black back")

Type: Constant current pulse

Mode: Continuous

Duration: 0.2 ms (200 microseconds)

Rate: 1.1/sec

Laterality: Unilateral (uninvolved side first)

Intensity: To produce supra-maximal response

(usually > 10 mA)





ENoG: Acquisition Parameters

Analysis time: 20 ms

Filter settings: 3 to 3000 Hz

Notch filter: No

Amplification: X 5000 or less

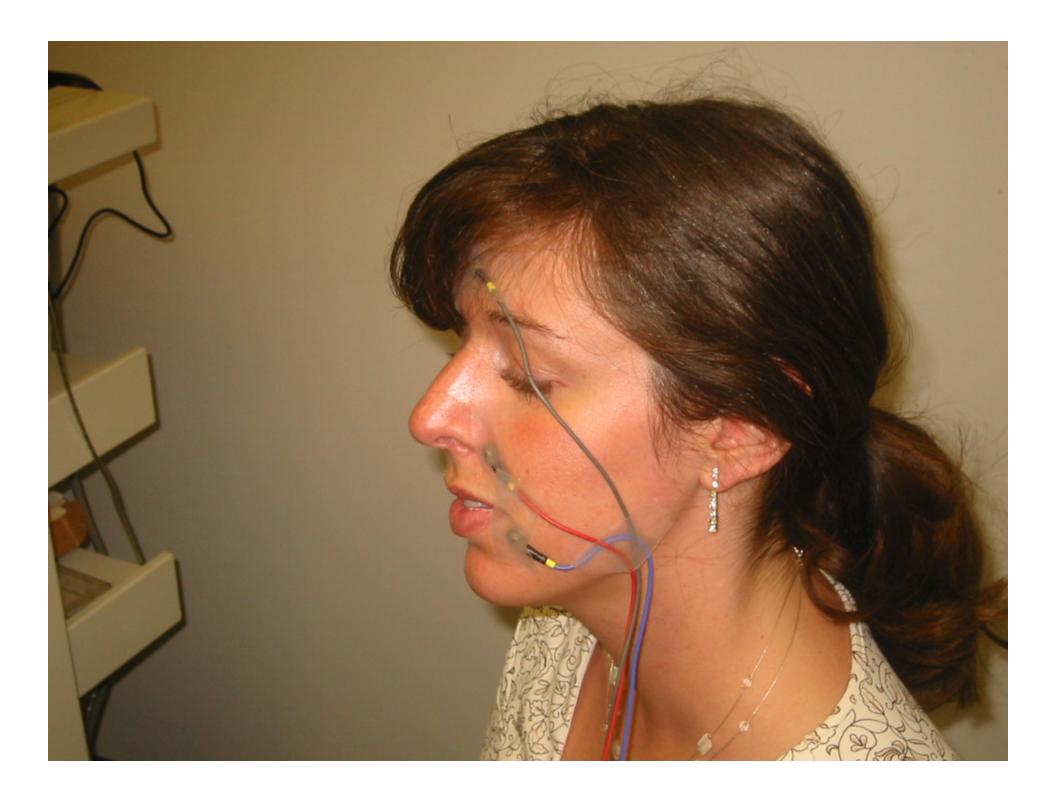
Electrodes: Ipsilateral nasolabial fold

(ground on forehead)

Impedance: < 5K ohms

Sweeps: 1 to 20





Electroneuronograpy (ENoG): Principles

Patients motivated to find a definitive diagnosis

They will tolerate some discomfort to reach the diagnosis.

Fully explain ENoG procedure, avoiding terms that will cause alarm or concern.

Possible explanations for a "sub-optimal" response should be explored and resolved with trouble-shooting and modification of the technique.



ENoG Principles (continued)...

Evoke optimal ("supra-maximal") distal facial nerve EMG from each side w/same stimulation and recording technique

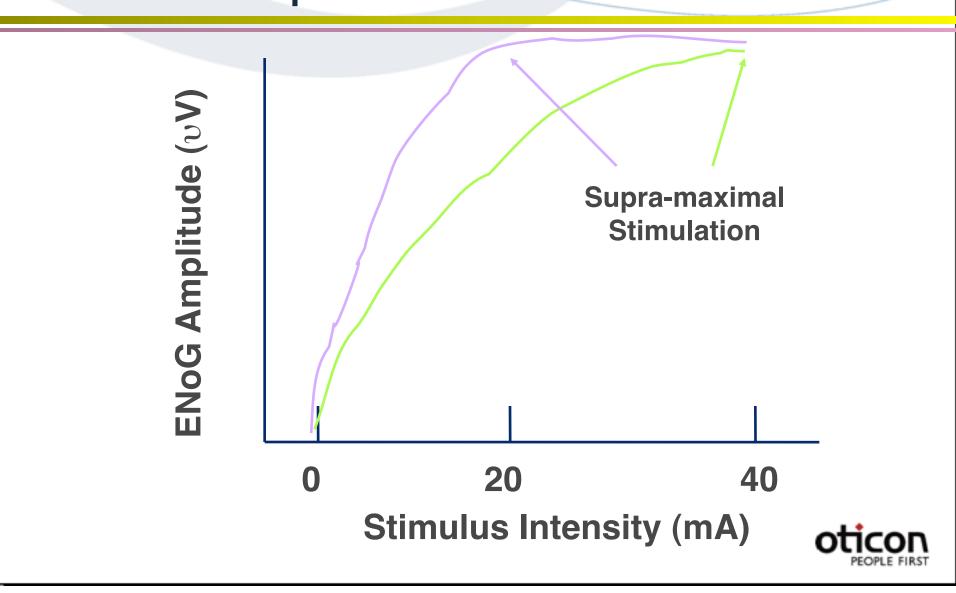
Analyze "bad" relative to "good" side.

Amplitude matters, not latency.

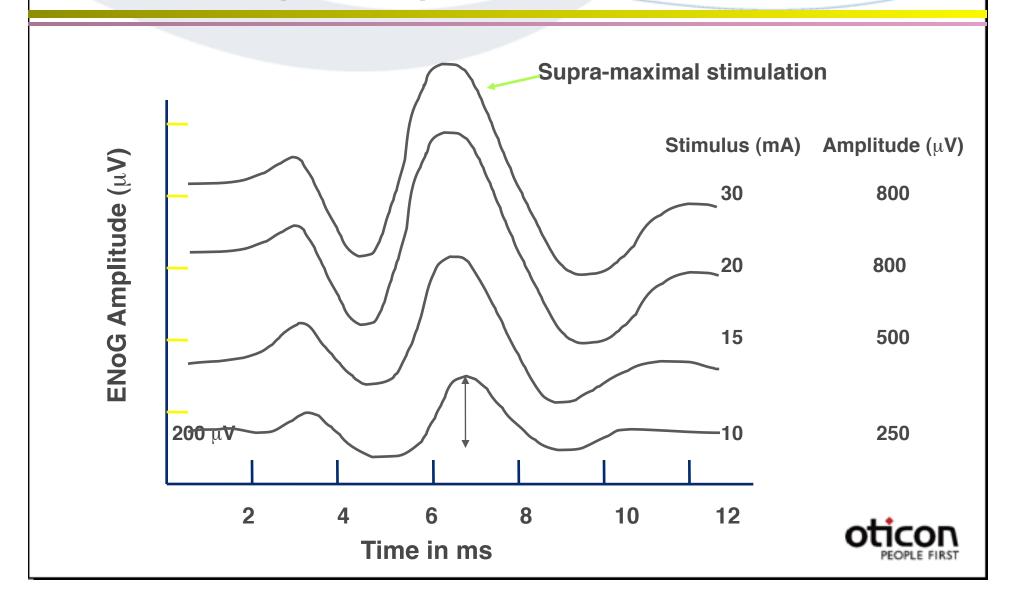
If "bad" side <10% of good side, significant degeneration has occurred, surgical intervention is an option.

If "bad" side >10%, possible spontaneous recovery ... "watch & wait".

ENoG Stimulus Parameters: Supramaximal Stimulation

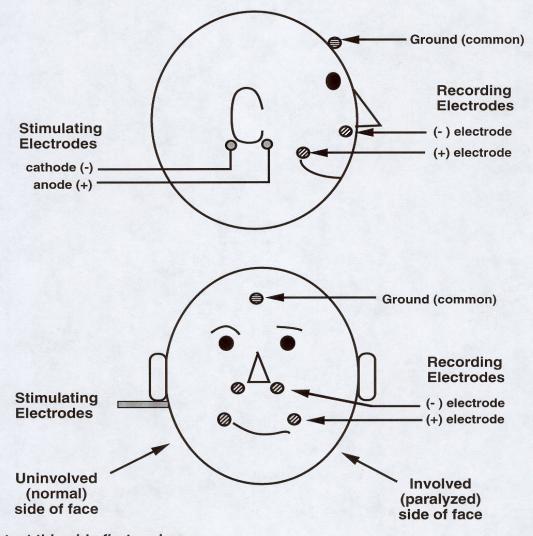


ENoG: Concept of Supra-Maximal Stimulation



ELECTRONEURONOGRAPHY (ENoG)

Stimulating and Recording Electrode Sites

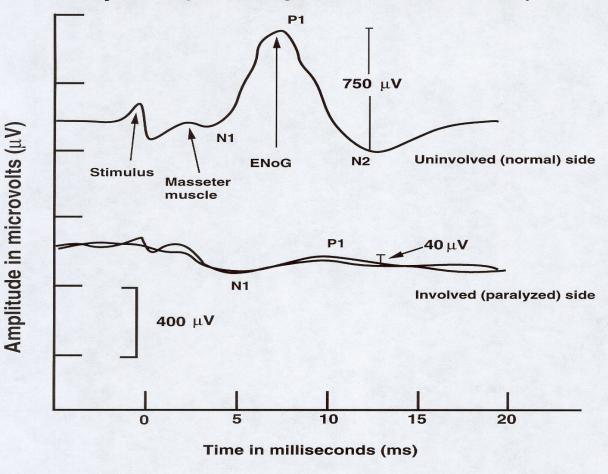


TIP: test this side first and use ENoG values as a reference



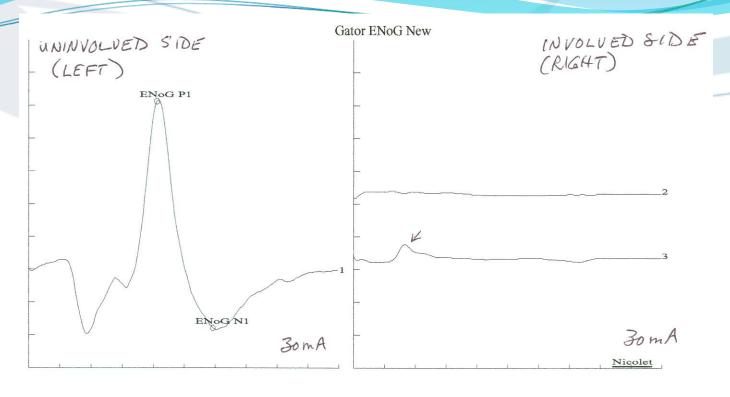
Electroneuronography (ENoG) Analysis:

Analysis: "Significant degeneration in facial nerve response"



Percent degeneration = 100 -
$$\frac{}{750~\mu\text{V}}$$
 X 100 = 95%





Sensitivity and Sweep Time Per Division

1 | 250.00 uV 2.0 msec | 2 | 250.00 uV 2.0 msec | 3 | 250.00 uV 2.0 msec

E	NoG	
12.84ms	ENoG Amp	
>12 IIII		



ENoG Recording:Trouble Shooting

Symptom: No response bilaterally

Problems: Chemical paralysis (in the O.R. or ICU)

Improper electrode placement

Electrical failure

Solutions: Reverse neuromuscular blocking agent medically

Verify correct electrode placement

Verify stimulation



Symptom: Poor response bilaterally

Problems: Edema at stimulation side (in trauma)

Tenderness/pain precludes adequate stimulus

electrode pressure

Solutions: Pain relief

Defer recording to later time



Symptom: Poor response bilaterally

Problems: Obese patient

Ineffective stimulation

Bilateral dysfunction or injury

Solutions: Apply pressure to stimulating electrodes

Needle electrodes

Compare patient ENoG amplitudes to normal data



Symptom: Excessive artifact rejection (cannot average)

Problems: Stimulus artifact

Very large normal response

Solutions: Increase distance btw stim & recording electrodes

Avoid crossing stim/rec electrode wires

Use post-stimulus delay

Decrease amplification (gain)



Symptom: Early response (peak before 6 ms)

Problem: Masseter muscle response

Solution: Move stimulating electrode posteriorly



Who introduced electroneuronography as a clinical procedure, and coined the term:

- a. Hallowell Davis
- b. James Jerger
- c. Robert Galambos
- d. Ugo Fisch
- e. Derrald Brackmann



Which of the following diseases is associated with facial paralysis?

- a. Herpes zoster
- b. Guillain Barre syndrome
- c. Temporal bone fracture
- d. Bell's palsy
- e. All of the above



The facial nerve exits the skull at the:

- a. Internal auditory canal
- b. Foramen magnum
- c. Stylomastoid foramen
- d. Facial canal
- e. Gerhardt's passageway



The facial nerve is a purely motor nerve:

- a. True
- b. False



The term describing total severing of the facial nerve is:

- a. Neuropraxia
- b. Neurotmesis
- c. Axontmesis
- d. Discontinuity
- e. Paralysis



Diabetics are more likely to develop Bell's palsy than non-diabetics:

- a. True
- b. False



The best time frame for ENoG measurement and clinical value is:

- a. Immediately upon injury
- b. Within 24 hours of onset of facial nerve dysfunction
- c. 72 hours after onset
- d. 21 days after onset
- e. Any of the above times are O.K.



The facial nerve exits the skull at the:

- a. Internal auditory canal
- b. Foramen magnum
- c. Stylomastoid foramen
- d. Facial canal
- e. Gerhardt's passageway



A typical stimulus in ENoG recording is:

- a. 80 dB HL
- b. 20 microvolts
- c. 20 amps
- d. 20 mA
- e. 50 mA



Grade 6 on the HB scale is:

- a. Normal facial nerve function
- b. No eye blink
- c. Ocular tremor
- d. Asymmetric smile
- e. Total facial paralysis



Which of the following pathological processes is measured in ENoG recording:

- a. Temporal bone fracture
- b. Bell's palsy
- c. Auditory-facial neuropathy (ANF)
- d. Wallerian degeneration
- e. Sensory facial nerve activation



The stimulating electrode and location is best described by which of the following:

- a. Anode anterior and cathode posterior
- b. Negative anterior and positive posterior
- c. Naso-labial fold
- d. Anode posterior and cathode anterior
- e. "Black" anterior and "red" posterior



The optimal stimulus intensity level for ENoG recording is:

- a. > 95 dB nHL
- b. > 40 mA
- c. 20 to 40 mA
- d. Supra-maximal intensity level
- e. Level producing facial nerve twitching



Recording electrodes in ENoG measurement are placed at the:

- a. Fz and SMF locations
- b. Stylomastoid foramen
- c. Forehead and corner of eye
- d. Nasolabial fold
- e. Corner of mouth and eye



A significant abnormality in ENoG recording is defined as:

- a. Amplitude of 20 microvolts
- b. Involved to non-involved ratio of 40%
- c. Degeneration of > 50% for involved vs. non-involved side
- d. Degeneration of > 90% for involved vs. non-involved side
- e. None of the above



A peak in an ENoG recording less than 6 ms is probably:

- a. PAM artifact
- b. Wave P1
- c. Sternocleidomastoid muscle
- d. Masseter muscle
- e. Buccal muscle

